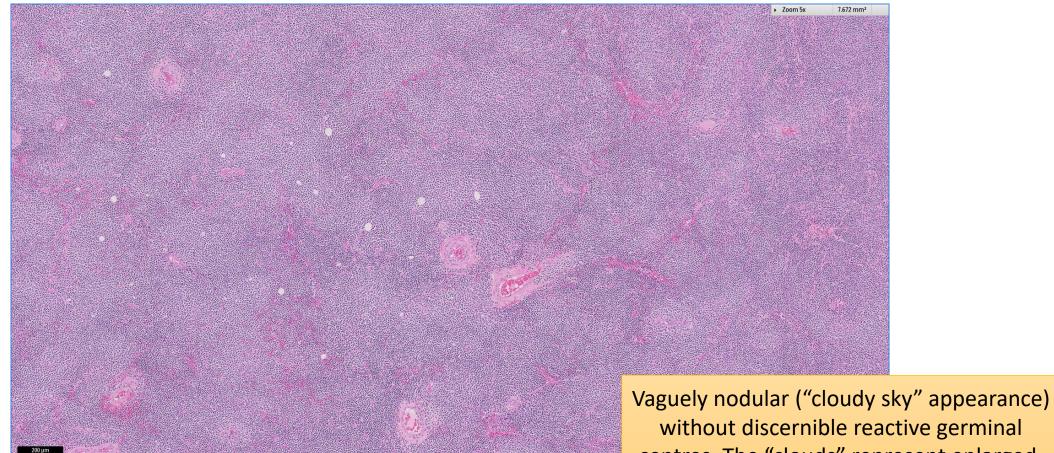
SGH CASE 26: 17RE144 & 17RE143

- Clinical History:
 - 60+ years old, Female. Hepatosplenomegaly and lymphadenopathy. Peripheral blood lymphocytosis (49.11 X 10^9/L) with prolymphocytes (5.78 x 10^9/L).
- Specimen:
 - Bone marrow trephine biopsy (17RE143) and lymph node biopsy (17RE144)

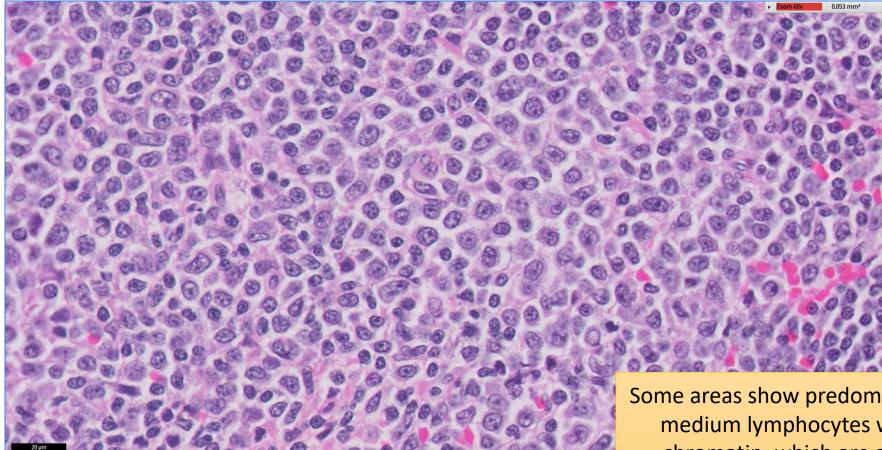
SGH CASE 26: 17RE144 LYMPH NODE H&E



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without discernible reactive germinal centres. The "clouds" represent enlarged, coalescent proliferation centres.

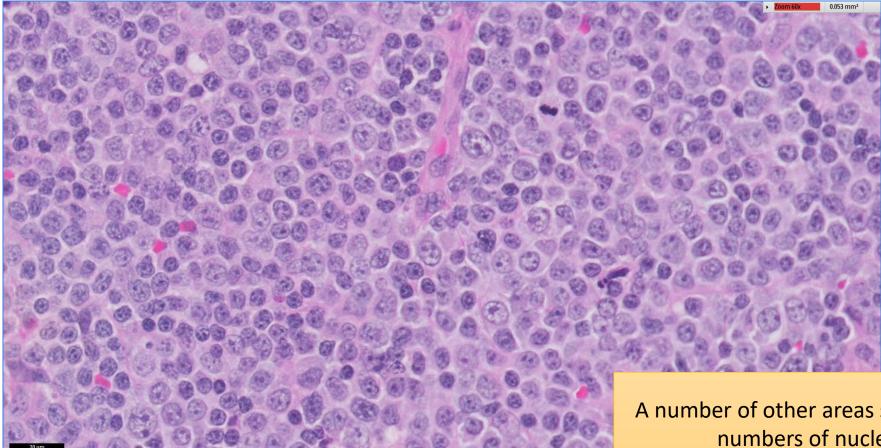
SGH CASE 26: 17RE144 LYMPH NODE H&E



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Some areas show predominantly small-tomedium lymphocytes with clumped chromatin, which are admixed with nucleolated cells

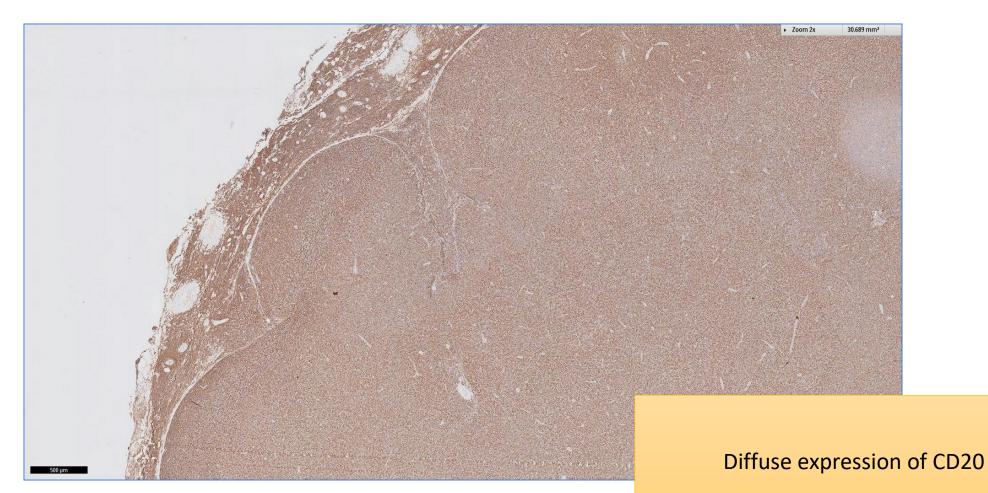
SGH CASE 26: 17RE144 LYMPH NODE H&E



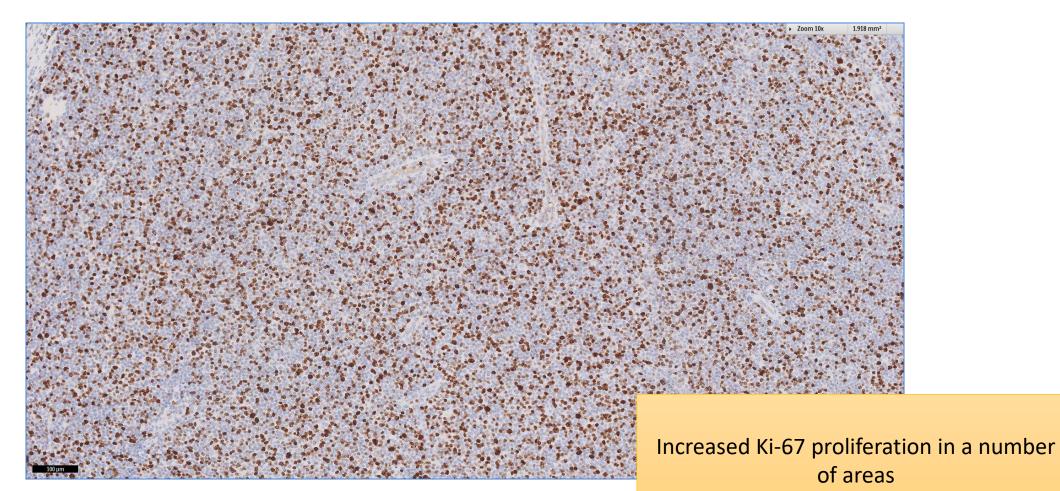
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A number of other areas show increased numbers of nucleolated paraimmunoblasts

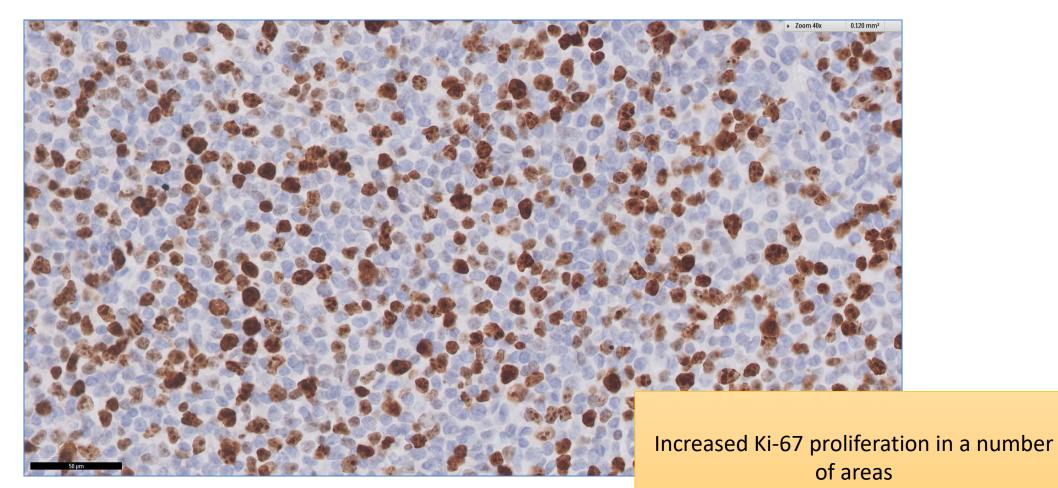
SGH CASE 26: 17RE144 LYMPH NODE CD20



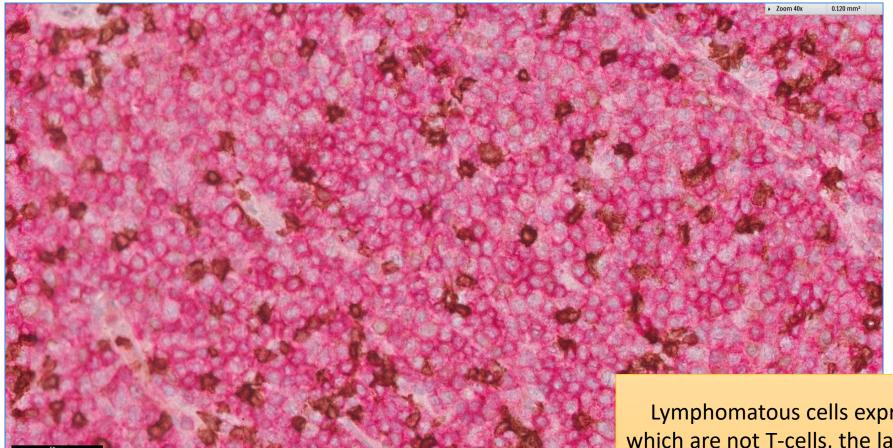
SGH CASE 26: 17RE144 LYMPH NODE KI67



SGH CASE 26: 17RE144 LYMPH NODE KI67

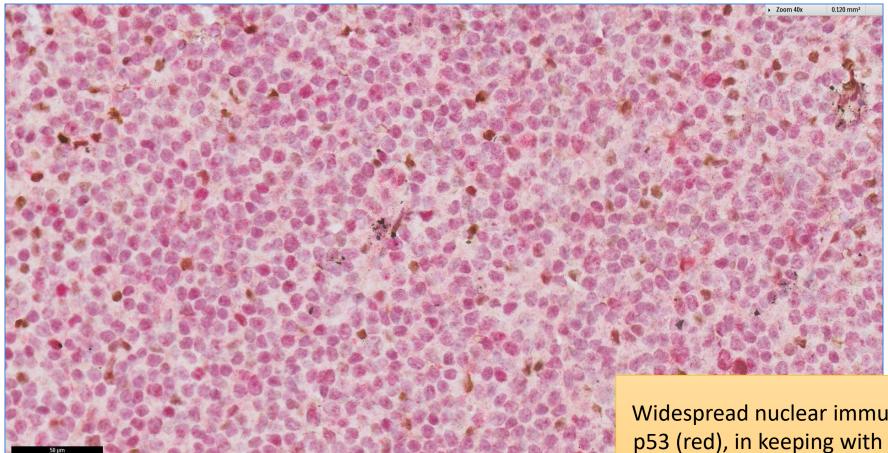


SGH CASE 26: 17RE144 LYMPH NODE CD5(RED) CD3(BROWN)



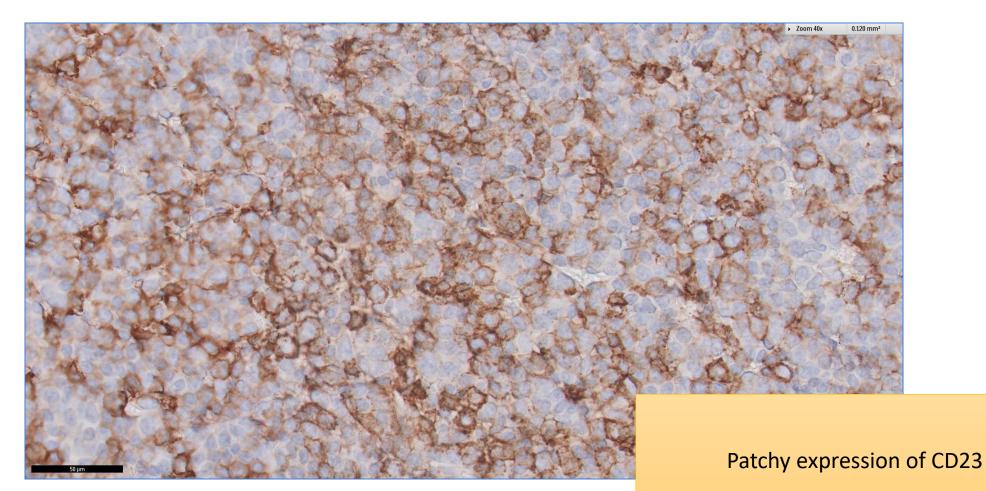
Lymphomatous cells express CD5(red) which are not T-cells, the latter highlighted by CD3

SGH CASE 26: 17RE144 LYMPH NODE P53(RED) P21(BROWN)

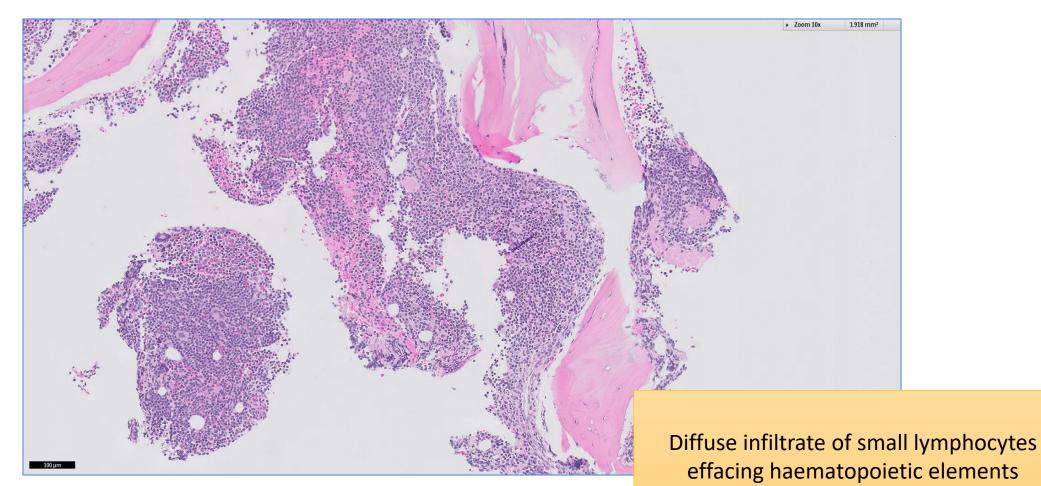


Widespread nuclear immunoreactivity for p53 (red), in keeping with mutant protein expression

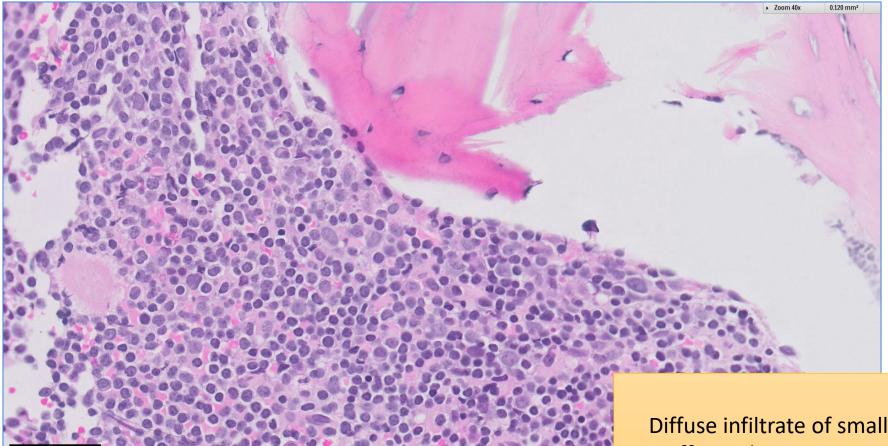
SGH CASE 26: 17RE144 LYMPH NODE CD23



SGH CASE 26: 17RE143 BONE MARROW H&E



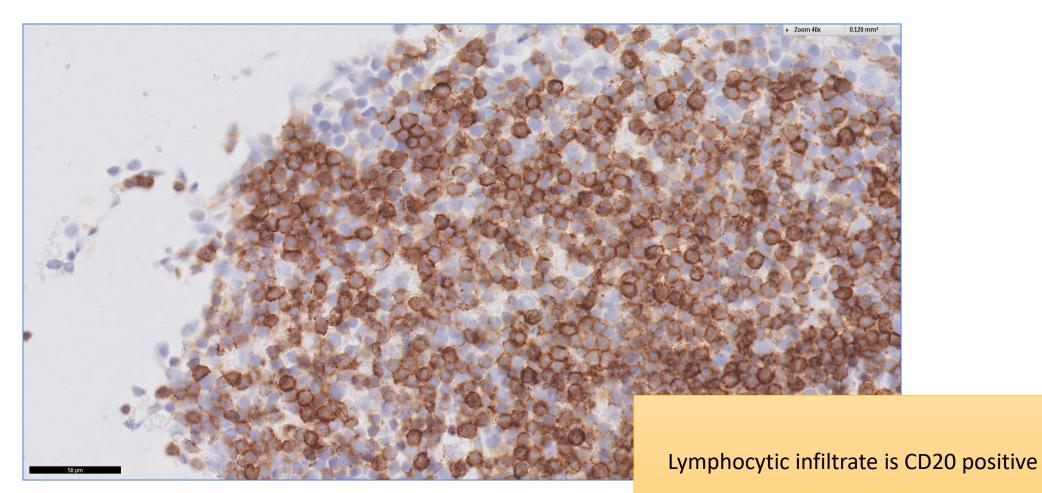
SGH CASE 26: 17RE143 BONE MARROW H&E



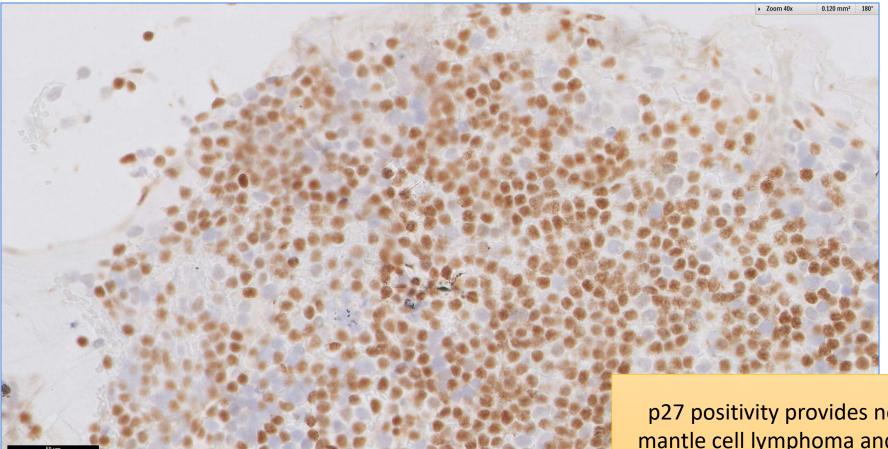
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Diffuse infiltrate of small lymphocytes effacing haematopoietic elements

SGH CASE 26: 17RE143 BONE MARROW CD20



SGH CASE 26: 17RE143 BONE MARROW P27



p27 positivity provides no support for mantle cell lymphoma and also indicate quiescence

SGH CASE 26: 17RE144 & 17RE143

- 17RE144 (Lymph node)
 - Chronic Lymphocytic Leukaemia In-transformation to Large B-cell Lymphoma (Richter Syndrome)
- 17RE143 (Bone marrow)
 - Involved by Chronic Lymphocytic Leukaemia